FDA 510(k) Premarket Notification

K955352 SUCTION CAUTERY PROBE

CLEARED

Apr 5, 1996
136 days to decision

K955352 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k955352/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 21, 1995
Decision date	Apr 5, 1996
Days to decision	136 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Mectra Labs, Inc.**
Location	Bloomfield, IN, US
Contact	CHARLES E ALLGOOD
510(k) history	9 submissions · 9 cleared · 1992-2012